

The listing of claims presented below replaces all prior versions and listing of claims in the application.

Listing of claims:

1. (Previously presented) A dosage form of a pharmaceutical composition which comprises an effective amount of a centrally-acting acetylcholinesterase inhibitor for the treatment of Alzheimer's disease selected from the group consisting of galanthamine, lycoramine, analogs of galanthamine, analogs of lycoramine and rivastigmine wherein said analogs of galanthamine or lycoramine are compounds wherein one or more of the methoxy, hydroxy or N-methyl groups is replaced as follows: the methoxy group by another alkoxy group of from two to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzyloxy or substituted benzyloxy group, a carbonate group or a carbamate group; the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzyloxy or substituted benzyloxy group, a carbonate group or a carbamate group; the N-methyl group by hydrogen, alkyl, benzyl or a cyclopropylmethyl group or a substituted or unsubstituted benzyloxy group; said compounds having a half life of from one to eleven hours wherein the acetylcholinesterase inhibitor is formulated so as to delay its activity for a predetermined period of from four to twelve hours such that acetyl-cholinesterase inhibition is avoided during such predetermined period.

2. (Canceled)

3. (Previously presented) A dosage form of a pharmaceutical composition as claimed in claim 1 wherein the composition is formulated to delay the activity of the acetyl cholinesterase inhibitor for a period of from six to nine hours.

4. (Previously presented) A dosage form of a pharmaceutical composition as claimed in claim 1 wherein the composition is formulated to delay the activity of the acetyl cholinesterase inhibitor for a period of from eight to twelve hours.

5. (Original) A dosage form of a pharmaceutical composition as claimed in claim 1 wherein said acetylcholinesterase inhibitor has a duration of action of from 2 to 12 hours.

6. (Canceled)

7. (Canceled)

8. (Canceled)

9. (Canceled)

10. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of galanthamine, lycoramine and analogs thereof wherein the methoxy group of such compounds is replaced by a hydrogen, hydroxy or alkoxy group of from two to six carbon atoms or an acyloxy group of from one to seven carbon atoms.

11. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine or lycoramine wherein the methoxy group thereof is replaced by a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.

12. (Previously presented) A dosage form as claimed in claim 11 wherein the alkyl group or groups of said carbonate or carbamate groups comprise from 4 to 6 carbon atoms.

13. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine or lycoramine wherein the hydroxy group thereof is replaced by a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.

14. (Previously presented) A dosage form as claimed in claim 12 wherein the alkyl group or groups of said carbonate or carbamate groups comprise from 4 to 6

carbon atoms.

15. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine or lycoramine wherein the methoxy group thereof is replaced by an aryl carbamate or carbonate group wherein said aryl group is selected from phenyl, naphthyl, substituted phenyl and substituted naphthyl groups wherein said substituent is selected from alkyl and alkoxy groups of from 1 to 6 carbon atoms, trifluoro methyl groups and halo groups.

16. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine and lycoramine wherein the hydroxy group thereof is replaced by an aryl carbamate or carbonate group wherein said aryl group is selected from phenyl, naphthyl, substituted phenyl and substituted naphthyl groups wherein said substituent is selected from alkyl and alkoxy groups of from 1 to 6 carbon atoms, trifluoro methyl groups and halo groups.

17. (Previously presented) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is selected from the group consisting of galanthamine, lycoramine and analogs thereof wherein the hydroxy group of such compounds is replaced by a hydrogen or alkoxy group of from one to six carbon atoms or an acyl group of from one to seven carbon atoms.

18. (Original) A dosage form of a pharmaceutical composition as claimed in claim 7 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine and lycoramine wherein the hydroxy group of galanthamine or lycoramine is replaced by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group.

19. (Original) A dosage form as claimed in claim 7 wherein said acetylcholinesterase inhibitor is galanthamine.

20. (Original) A dosage form as claimed in claim 1 wherein said acetylcholinesterase inhibitor is rivastigmine.

21. (Canceled)

22. (Canceled)

23. (Canceled)

24. (Previously presented) A method of treatment as claimed in claim 41 wherein the composition is formulated to delay release of the acetyl cholinesterase inhibitor for a period of from six to nine hours.

25. (Previously presented) A method of treatment as claimed in claim 41 wherein the composition is formulated to delay release of the acetyl cholinesterase inhibitor for a period of from eight to twelve hours.

26. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor has a duration of action of from 2 to 12 hours.

27. (Canceled)

28. (Canceled)

29. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is selected from the group consisting of galanthamine, lycoramine and analogs thereof wherein the methoxy group of such compounds is replaced by a hydrogen, hydroxy or alkoxy group of from two to six carbon atoms or an acyloxy group of from one to seven carbon atoms.

30. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of

galanthamine or lycoramine wherein the methoxy group thereof is replaced by a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.

31. (Original) A method of treatment as claimed in claim 30 wherein the alkyl group or groups of said carbonate or carbamate groups comprise from 4 to 6 carbon atoms.

32. (Original) A method of treatment as claimed in claim 31 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine or lycoramine wherein the hydroxy group thereof is replaced by a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.

33. (Original) A method of treatment as claimed in claim 32 wherein the alkyl group or groups of said carbonate or carbamate groups comprise from 4 to 6 carbon atoms.

34. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine or lycoramine wherein the methoxy group thereof is replaced by an aryl carbamate or carbonate group wherein said aryl group is selected from phenyl, naphthyl, substituted phenyl and substituted naphthyl groups wherein said substituent is selected from alkyl and alkoxy groups of from 1 to 6 carbon atoms, trifluoro methyl groups and halo groups.

35. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is selected from the group consisting of analogs of galanthamine and lycoramine wherein the hydroxy group thereof is replaced by an aryl carbamate or carbonate group wherein said aryl group is selected from phenyl, naphthyl, substituted phenyl and substituted naphthyl groups wherein said substituent is selected from alkyl and alkoxy groups of from 1 to 6 carbon atoms, trifluoro methyl groups and halo groups.

36. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is selected from the group consisting of galanthamine, lycoramine and analogs thereof wherein the hydroxyl group of such compounds is replaced by a hydrogen or alkoxy group of from one to six carbon atoms or an acyl group of from one to seven carbon atoms.

37. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is galanthamine.

38. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is rivastigmine.

39. (Previously presented) A method of treatment as claimed in claim 41 wherein said acetylcholinesterase inhibitor is administered in conjunction with a compound that reduces its peripheral effects.

40. (Previously presented) A method of treatment as claimed in claim 39 wherein said acetylcholinesterase inhibitor is administered in conjunction with a suitable dose of propantheline bromide or glycopyrrolate.

41. (Previously presented) A method of treatment of a patient suffering from Alzheimer's disease which comprises administering a delayed release dosage form of a pharmaceutical composition which comprises an effective amount for treating Alzheimer's disease of a centrally acting acetylcholinesterase inhibitor selected from the group consisting of galanthamine, lycoramine, analogs of galanthamine and analogs of lycoramine and rivastigmine wherein said analogs of galanthamine or lycoramine are compounds wherein one or more of the methoxy, hydroxy or N-methyl groups is replaced as follows: the methoxy group by another alkoxy group of from two to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a

carbonate group or a carbamate group; the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group; the N-methyl group by hydrogen, alkyl, benzyl or a cyclopropylmethyl group or a substituted or unsubstituted benzoyloxy group; wherein said dosage is formulated such that the half life of the activity of the acetylcholinesterase inhibitor and the degree of delayed release are selected such that the formulation may be administered to the patient such that release of acetylcholinesterase inhibitor is avoided for the next anticipated sleep time and wherein administration of the drug is at an appropriate time to achieve such results.

42. (Previously presented) A method of treating Alzheimer's disease comprising administration to a patient suffering therefrom a composition comprising a centrally acting acetyl-cholinesterase inhibitor selected from the group consisting of galanthamine, lycoramine, analogs of galanthamine, analogs of lycoramine and rivastigmine wherein said analogs of galanthamine or lycoramine are compounds wherein one or more of the methoxy, hydroxy or N-methyl groups is replaced as follows: the methoxy group by another alkoxy group of from two to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group; the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group; the N-methyl group by hydrogen, alkyl, benzyl or a cyclopropylmethyl group or a substituted or unsubstituted benzoyloxy group; said compounds having a half life of from one to eleven hours wherein the nature of the composition and the time of administration of said acetyl cholinesterase inhibitor from said composition is such as to minimize release prior and during to desired sleep hours so as to allow the patient's central nervous system to become hypocholinergic during the period of desired sleep as compared to its cholinergic activity during hours when desired to be awake SO as to superimpose a physiological rhythm of cholinergic activity to a brain in which the cholinergic system is deteriorating.